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**CENTRAL FAX CENTER** Appl. No. 10/805,099 (Docket 099/004)  
**DEC 13 2006** Arndt, dated Dec. 13, 2006  
Reply to Office Action of July 13, 2006

**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

What is claimed as the invention is:

The invention claimed is:

1. A method of generating a cell composition containing cardiomyocytes or cardiomyocyte precursor cells from ~~primate pluripotent stem (pPS) cells obtained from a human blastocyst or human embryonic stem (hES) cells~~, comprising:
  - a) initiating differentiation of the pPS hES cells in suspension culture by forming embryoid bodies;
  - b) culturing the initiated cells so that they differentiate ~~into areas that undergo spontaneous contraction~~;
  - c) harvesting the differentiated cells;
  - d) separating the harvested cells into fractions according to their based on density; and
  - e) collecting combining the cell fractions containing cells that express cardiac troponin I (cTnI), cardiac troponin T (cTnT), or atrial natriuretic factor (ANF) from an endogenous gene;  
thereby generating a cell composition containing cardiomyocytes or cardiomyocyte precursor cells.
2. The method of claim 1, wherein the embryoid bodies are plated onto a surface coated with gelatin or Matrigel®.
3. The method of claim 1, wherein the cells are differentiated in the presence of a nucleotide analog that affects DNA methylation, such as 5-aza-deoxy-cytidine.

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4. The method of claim 1, wherein the cells are differentiated in a growth environment comprising a morphogen such as activin, and two or more growth factors.
5. The method of claim 4, wherein the morphogen is an activin, and the growth factors include an insulin-like growth factor and a member of the TGF $\beta$  family.
6. The method of claim 1, wherein the cells are differentiated in a growth environment containing about 20% serum or serum substitute.
7. The method of claim 1, wherein the harvested cells are separated by density centrifugation.
8. The method of claim 1, wherein the separating comprises distributing cells in the population according to based on their density, and collecting cells at combining cell fractions with a density between ~1.05 and ~1.075 g/mL.
9. The method of claim 1, further comprising culturing the collected cells combined cell fractions for at least 1 week in a medium containing a compound capable of forming a high energy phosphate bond, an acyl group carrier molecule, and a cardiomyocyte calcium channel modulator.
10. The method of claim 9, further comprising culturing the collected cells combined cell fractions for at least 1 week in a medium containing creatine, carnitine, or taurine.
- 11.-16. (Canceled)